Assessment of the genetic diversity of wild kuruma prawn based on the relatedness analysis using microsatellites DNA and mitochondrial DNA markers

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Huge amounts of the prawns have been released around Japan

15 - 20 millions/year

Japan Sea

50 millions/year

Seto Inland Sea

40 millions/year

East China Sea

35 - 40 millions/year

Pacific Ocean



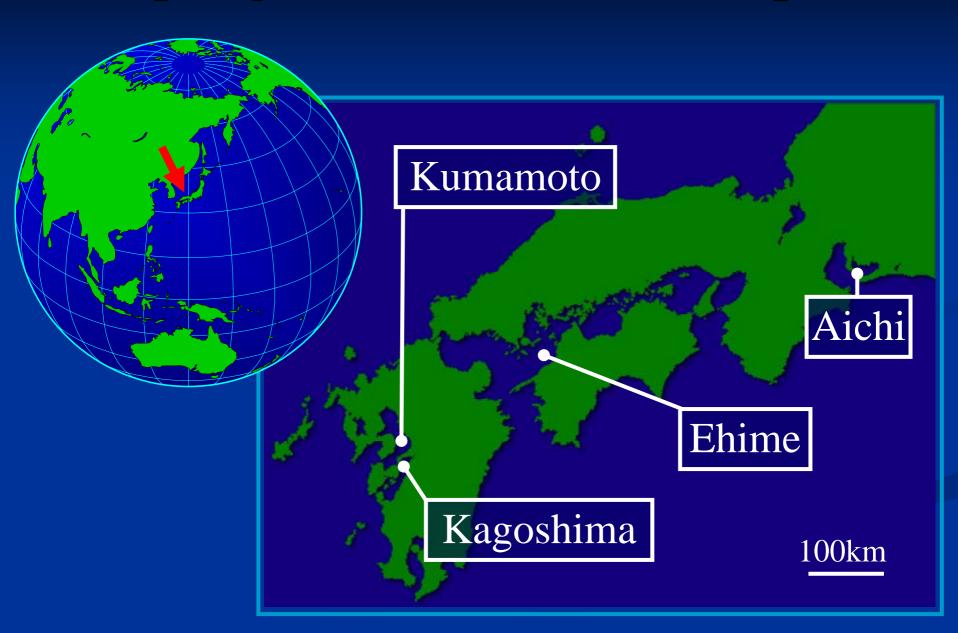
...However, cost and benefit of the stocking are still ambiguous.

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 - ·Population genetic analyses based on allele frequency
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 - ·Population genetic analyses based on relatedness

Sampling locations of kuruma prawn



The DNA markers used in this study

Microsatellites DNA (MS-DNA) marker

Loci	Repeat motif			
<i>CSPJ002</i> *	$(GA)_n$			
<i>CSPJ010</i> *	$(GAA)_nGCA(GAA)_m$			
<i>CSPJ012</i> *	$(CTT)_nCCT(CTT)_mCAT(CTT)_1$			
<i>CSPJ014</i> *	GAAAA(GAA) _n GAAAA			
<i>CSPJ015</i> *	$(CTT)_n$			

Mitochondrial DNA (mtDNA) PCR-RFLP marker



The heterogeneities of prawns in Kumamoto and Kagoshima were suggested by HWE test

Loci	Aichi	Ehime	Kuma- moto	Kago- shima
CSPJ002*	-	_		
<i>CSPJ010</i> *		_		
<i>CSPJ012</i> *	-	- (_	
<i>CSPJ014</i> *				
<i>CSPJ015</i> *	-		_	_

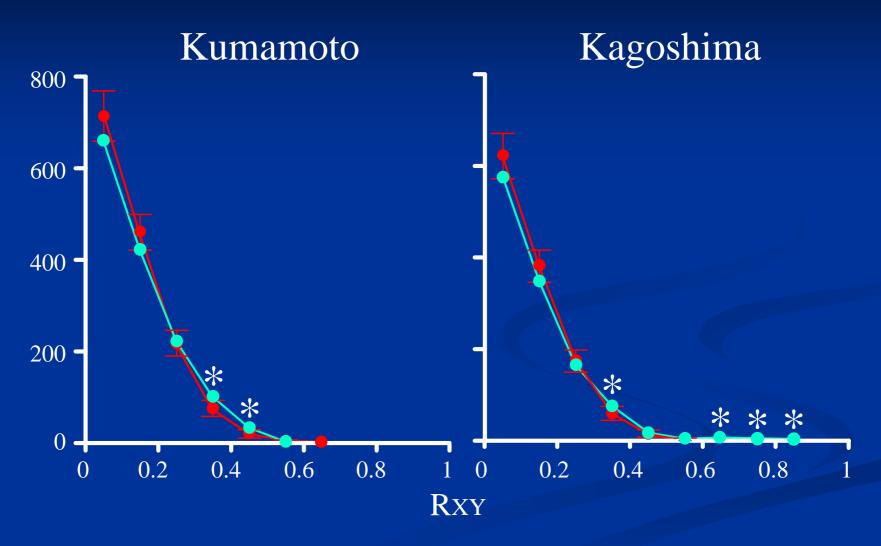
+: Significantly deviated from Hardy-Weinberg equilibrium (P<0.05)

Significant differences of haplotype distributions of the mtDNA marker were detected among the localities

	Aichi	Ehime	Kuma- moto
Ehime	_		
Kumamoto	_		
Kagoshima	+	+	+

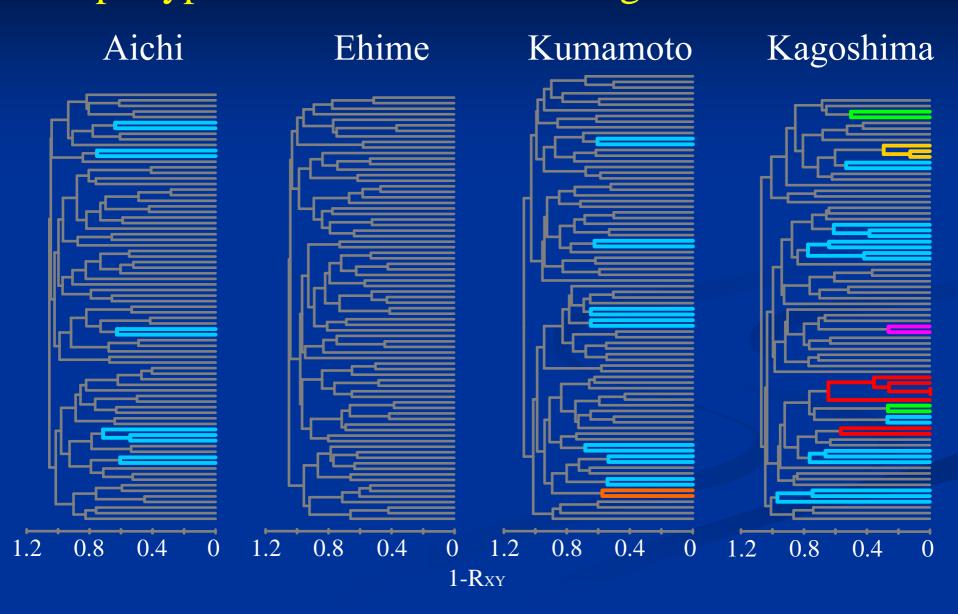
+: Significant (*P*<0.05)

Rxy tended to be high in Kumamoto and Kagoshima



•:Observed value •:Simulated value *:Significant (P<0.05)

Closely related individuals tended to share mtDNA haplotypes in Kumamoto and Kagoshima



Conclusion

Genetic analyses using MS-DNA markers

- · Genetic diversity were high in all localities.
- · HWE tests suggested heterogeneity within the localities.
- → Kin groups should be existed in the localities.
- · No genetic difference was detected among the localities.
- → Gene flows among the localities are large.

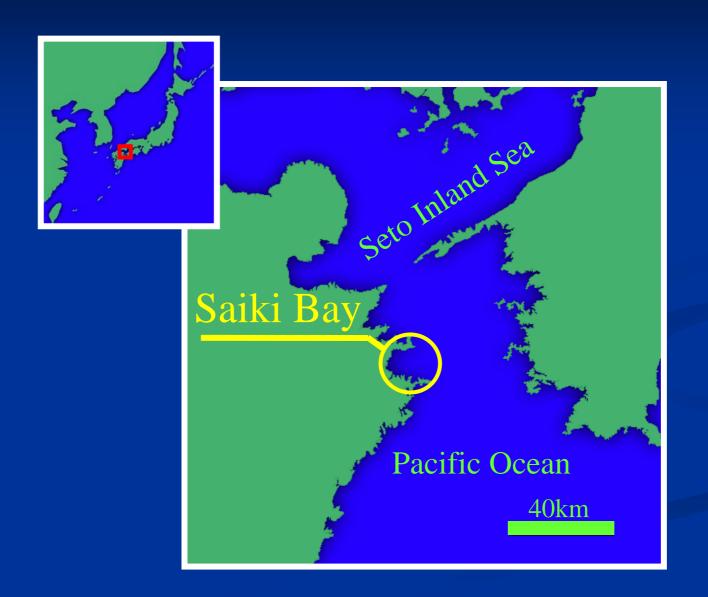
Genetic analyses using mtDNA markers

- · Genetic diversity were high in all localities.
- · Genetic differences were significant among the localities.
- → It is caused by the heterogeneity within the localities.

The relatedness should be useful for the detailed population genetic analysis.

Assessment of genetic diversity of stocked population

Location of the survey area in this analysis



Detail of the prawn stocking at Saiki Bay in 2005

No. of released prawns: 693,000

No. of marked prawns: 197,000

Size at release: 5cm in TL

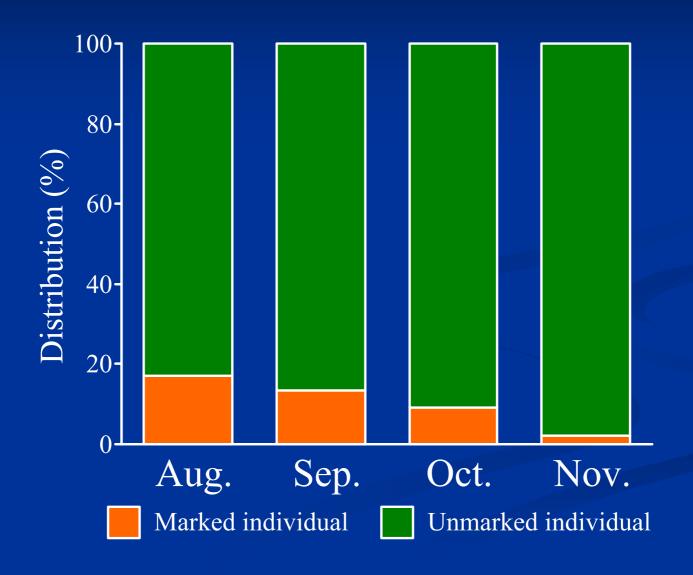
Date of release: from Jul. to Aug. '05

Marked prawn

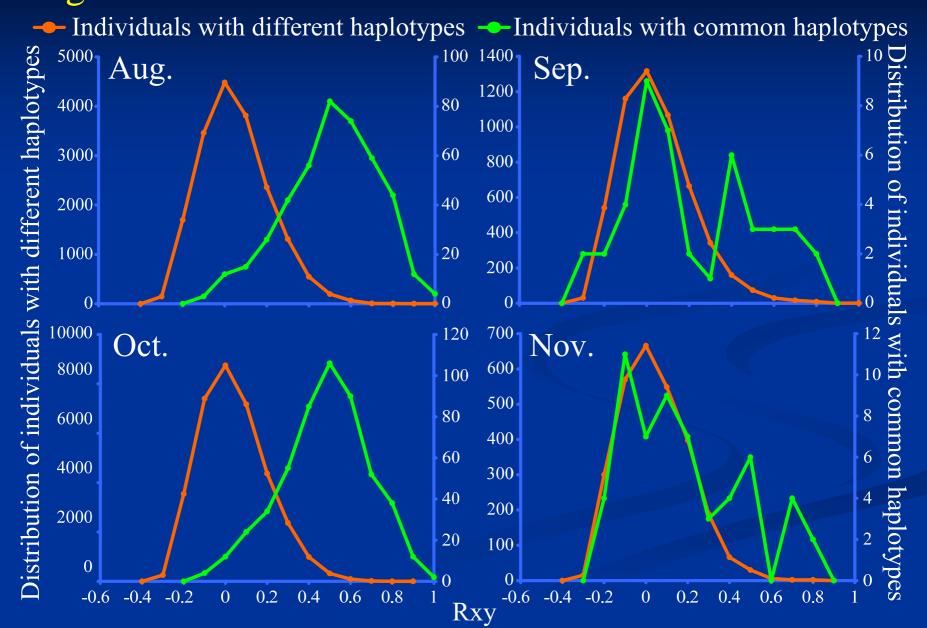
A right side of caudal appendage was cut off



Distributions of marked individuals in the prawns caught in Saiki Bay

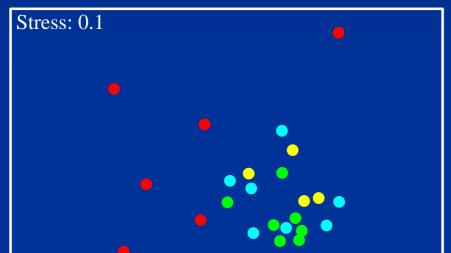


The individuals sharing haplotypes showed relatively high relatedness

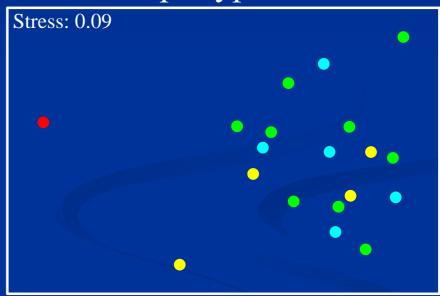


Genetic relatedness among individuals with common haplotypes

Haplotype #3



Haplotype #30



Month of sampling



Aug. Sep. Oct. Nov.

Future directions

- 1. Assessment of genetic diversity of the wild population
 - · Investigation of the degree of gene flow among localities.
 - · Investigation of population dynamics in each localities.
- 2. Assessment of genetic diversity of stocked population
 - · Investigation of the stocking effectiveness.
 - · Investigation of the genetic influences of the stocking.